SCOUT Accessories for Implanting in the Coronary Sinus 510(k) K020821

1. 510(K) SUMMARY

Name and Address of Sponsor: BIOTRONIK, Inc.
6024 Jean Road
Lake Oswego, OR 97035

Establishment Registration Number: 1028232

Device Name:

Proprietary Name:	SCOUT
Classification:	Class II (21 CFR 870.1330; 21 CFR 870.1380)
Classification Name:	Wire, Guide, Catheter; Stylet, Catheters
Product Code:	DQX, DRB

General Description:

The SCOUT Accessories for Implanting in the Coronary Sinus are for introducing legally marketed leads into the vessels of the left heart via the coronary sinus. The following SCOUT Accessories for Implanting in the Coronary Sinus are the subject of this 510(k):

- 1 - Hemostatic Valve
- 2 - Guiding Catheters Peel-Away (Long Sheaths)
- 1 - Dilator (for Long Sheath)
- 1 - Introducer Peel-Away 11F
- 1 - Guidewire
- 1 - Cannula
- 1 - Syringe Body
- 1 - Peel Tool

The SCOUT Accessories for Implanting in the Coronary Sinus are functionally equivalent to market-released delivery systems. In specific, the materials used and design are similar to Medtronic's LDS 6216 Left-Heart Delivery System (K012130, cleared 08-28-01).

Indication for Use:

The intended use of the SCOUT Accessories for Implantation in the Coronary Sinus is for introducing legally marketed leads into the vessels of the left heart via the coronary sinus.

Name and Address of Manufacturer:

BIOTRONIK GmbH & Co. (reg. no. 9610139)
Woermannkehre 1, 12359 Berlin, Germany
011-49-30-689-05-304

Contact Person(s) and Phone Number:

Jon Brumbaugh
Director, Regulatory Affairs
Phone (888) 345-0374 Fax (503) 635-9936

Name and Address of Contract Manufacturing Site:

BIOTRONIK AG (reg. no. 8043892)
Ackerstrasse 6
8180 Bülach, Switzerland
011-41-1-864-5169

2. INDICATIONS FOR USE

The intended use of the SCOUT Accessories for Implanting in the Coronary Sinus is for introducing legally marketed leads into the vessels of the left heart via the coronary sinus.

See Appendix 1 for the 510(k) Indications for Use Form.
Device description and substantial

3. PROPOSED LABELING

3.1 TECHNICAL MANUAL

The Technical Manual for the SCOUT Accessories is provided in Appendix 2.

DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

JUN 0 3 2002

BIOTRONIK, Inc.
c/o Mr. Jon Brumbaugh
Director, Regulatory Affairs
6024 Jean Road
Lake Oswego, OR 97035

Re: K020821
Trade Name: SCOUT
Regulation Number: 21 CFR 870.1330, 870.1380, and 870.1250
Regulation Name: Wire, Guide, Catheter; Stylet, Catheter; Catheter, Percutaneous
Regulatory Class: Class II (two)
Product Code: DQX, DRB, and DQY
Dated: March 8, 2002
Received: March 13, 2002

Dear Mr. Brumbaugh:

We have reviewed your Section 510(k) premarket notification of intent to market the device referenced above and have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act) that do not require approval of a premarket approval application (PMA). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration.

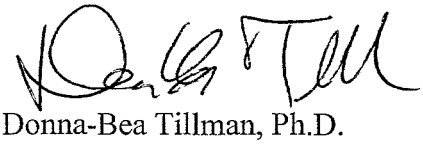
If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to such additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 898. In addition, FDA may publish further announcements concerning your device in the Federal Register.

Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies. You must comply with all the Act's requirements, including, but not limited to: registration and listing (21 CFR Part 807); labeling (21 CFR Part 801); good manufacturing practice requirements as set forth in the quality systems (QS) regulation (21 CFR Part 820); and if applicable, the electronic product radiation control provisions (Sections 531-542 of the Act); 21 CFR 1000-1050.

This letter will allow you to begin marketing your device as described in your Section 510(k) premarket notification. The FDA finding of substantial equivalence of your device to a legally marketed predicate device results in a classification for your device and thus, permits your device to proceed to the market.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801 and additionally 21 CFR Part 809.10 for in vitro diagnostic devices), please contact the Office of Compliance at (301) 594-4646. Additionally, for questions on the promotion and advertising of your device, please contact the Office of Compliance at (301) 594-4639. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21CFR Part 807.97). Other general information on your responsibilities under the Act may be obtained from the Division of Small Manufacturers, International and Consumer Assistance at its toll-free number (800) 638-2041 or (301) 443-6597 or at its Internet address http://www.fda.gov/cdrh/dsma/dsmamain.html

Sincerely yours,



Donna-Bea Tillman, Ph.D.
Acting Director
Division of Cardiovascular
and Respiratory Devices
Office of Device Evaluation
Center for Devices and
Radiological Health

Enclosure

K020821
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510(k) Number (if known): K020821

Device Name: SCOUT

Indications For Use:

The SCOUT Accessories for Implanting in the Coronary Sinus is for introducing legally marketed leads into the vessels of the left heart via the coronary sinus.

(PLEASE DO NOT WRITE BELOW THIS LINE-CONTINUE ON ANOTHER PAGE IF NEEDED)

Concurrence of CDRH, Office of Device Evaluation (ODE)

Division of Cardiovascular & Respiratory Devices
510(k) Number K020821

(Optional Format 3-10-98)

Prescription Use X